



University of Kentucky
UKnowledge

Physics and Astronomy Faculty Publications

Physics and Astronomy

1-12-2017

Doping Evolution of Magnetic Order and Magnetic Excitations in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$

Xingye Lu

Paul Scherrer Institut, Switzerland

D. E. McNally

Paul Scherrer Institut, Switzerland

M. Moretti Sala

European Synchrotron Radiation Facility, France

Jsaminka Terzic

University of Kentucky, jasminka.terzic@uky.edu

M. H. Upton

Argonne National Laboratory

See next page for additional authors

Right click to open a feedback form in a new tab to let us know how this document benefits you.

Follow this and additional works at: https://uknowledge.uky.edu/physastron_facpub

 Part of the [Condensed Matter Physics Commons](#)

Repository Citation

Lu, Xingye; McNally, D. E.; Sala, M. Moretti; Terzic, Jsaminka; Upton, M. H.; Casa, D.; Ingold, G.; Cao, Gang; and Schmitt, T., "Doping Evolution of Magnetic Order and Magnetic Excitations in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ " (2017). *Physics and Astronomy Faculty Publications*. 532.

https://uknowledge.uky.edu/physastron_facpub/532

This Article is brought to you for free and open access by the Physics and Astronomy at UKnowledge. It has been accepted for inclusion in Physics and Astronomy Faculty Publications by an authorized administrator of UKnowledge. For more information, please contact UKnowledge@lsv.uky.edu.

Authors

Xingye Lu, D. E. McNally, M. Moretti Sala, Jsaminka Terzic, M. H. Upton, D. Casa, G. Ingold, Gang Cao, and T. Schmitt

Doping Evolution of Magnetic Order and Magnetic Excitations in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ **Notes/Citation Information**

Published in *Physical Review Letters*, v. 118, issue 2, 027202, p. 1-6.

© 2017 American Physical Society

The copyright holder has granted permission for posting the article here.

Digital Object Identifier (DOI)

<https://doi.org/10.1103/PhysRevLett.118.027202>

Doping Evolution of Magnetic Order and Magnetic Excitations in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$

Xingye Lu,^{1,*} D. E. McNally,¹ M. Moretti Sala,² J. Terzic,^{3,4} M. H. Upton,⁵ D. Casa,⁵ G. Ingold,^{1,6}
G. Cao,^{3,4} and T. Schmitt^{1,†}

¹Research Department Synchrotron Radiation and Nanotechnology, Paul Scherrer Institut, CH-5232 Villigen PSI, Switzerland

²European Synchrotron Radiation Facility, BP 220, F-38043 Grenoble Cedex, France

³Department of Physics and Astronomy, University of Kentucky, Lexington, Kentucky 40506, USA

⁴Department of Physics, University of Colorado at Boulder, Boulder, Colorado 80309, USA

⁵Advanced Photon Source, Argonne National Laboratory, Argonne, Illinois 60439, USA

⁶SwissFEL, Paul Scherrer Institut, CH-5232 Villigen PSI, Switzerland

(Received 28 August 2016; revised manuscript received 25 October 2016; published 12 January 2017)

We use resonant elastic and inelastic x-ray scattering at the Ir- L_3 edge to study the doping-dependent magnetic order, magnetic excitations, and spin-orbit excitons in the electron-doped bilayer iridate $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ ($0 \leq x \leq 0.065$). With increasing doping x , the three-dimensional long range antiferromagnetic order is gradually suppressed and evolves into a three-dimensional short range order across the insulator-to-metal transition from $x = 0$ to 0.05, followed by a transition to two-dimensional short range order between $x = 0.05$ and 0.065. Because of the interactions between the $J_{\text{eff}} = \frac{1}{2}$ pseudospins and the emergent itinerant electrons, magnetic excitations undergo damping, anisotropic softening, and gap collapse, accompanied by weakly doping-dependent spin-orbit excitons. Therefore, we conclude that electron doping suppresses the magnetic anisotropy and interlayer couplings and drives $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ into a correlated metallic state with two-dimensional short range antiferromagnetic order. Strong antiferromagnetic fluctuations of the $J_{\text{eff}} = \frac{1}{2}$ moments persist deep in this correlated metallic state, with the magnon gap strongly suppressed.

DOI: 10.1103/PhysRevLett.118.027202

The layered Ruddlesden-Popper iridate series $\text{Sr}_{n+1}\text{Ir}_n\text{O}_{3n+1}$ ($n = 1, 2$) that hosts novel $J_{\text{eff}} = \frac{1}{2}$ Mott insulating states have recently attracted much interest owing to their potential for exploring novel collective quantum states by charge-carrier doping in the strong spin-orbit coupling (SOC) limit [1–8]. Distinct from 3d Mott insulators where strong on-site Coulomb electron correlation (U) dominates [9], the $J_{\text{eff}} = \frac{1}{2}$ Mott state in iridates is induced by cooperative interplay between crystal-field, SOC (~ 0.4 eV) and an intermediate U [4]. The novel Mott insulator Sr_2IrO_4 is similar to La_2CuO_4 in magnetic order, spin dynamics, and electronic structure [3–14]. It has been suggested that electron-doped Sr_2IrO_4 is an analogous system to hole-doped La_2CuO_4 [15,16], which is supported by numerous non-trivial experimental observations such as Fermi arcs, pseudogaps, d -wave gaps, and persistent paramagnons [17–24].

Compared with Sr_2IrO_4 , the bilayer $\text{Sr}_3\text{Ir}_2\text{O}_7$ bearing strong interlayer coupling is similar to bilayer cuprates [25] and shows unique advantages in exploring novel phases via electron doping since it retains the $J_{\text{eff}} = \frac{1}{2}$ Mott state while lying close to an insulator-to-metal transition (IMT) [7,8]. Because of its bilayer structure, $\text{Sr}_3\text{Ir}_2\text{O}_7$ exhibits a much smaller gap $\Delta E \approx 130$ meV than Sr_2IrO_4 ($\Delta E \approx 600$ meV) [7,26,27]. The strong interlayer coupling and magnetic anisotropy induce a c -axis G -type antiferromagnetic order

(AFM) below $T_N \approx 285$ K [8,28] and distinct magnons bearing a large magnon gap ≈ 90 meV [29,30]. Because of the small charge gap of $\text{Sr}_3\text{Ir}_2\text{O}_7$, one can expect a homogeneous metallic state to develop by minor charge carrier doping. Indeed, an IMT and a robust metallic state have been realized in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ for $x \gtrsim 0.05$ [31,32]. Since minor La dopants have little structural effect on IrO_2 layers while driving the system across the IMT, $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ provides an ideal platform for exploring novel phenomena arising from charge carrier doped $J_{\text{eff}} = \frac{1}{2}$ Mott states, in the presence of strong interlayer coupling and magnetic anisotropy [29,30]. However, the doping evolution of the ground state of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ is still under intense debate [32–35]. Angle-resolved photoemission (ARPES) measurements revealed a strong coherent quasiparticle peak and suggested a conventional, weakly correlated Fermi liquid ground state for $x > 0.05$ in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ [33]. On the other hand, a comprehensive study [32] and later ARPES measurements [35] on $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ suggested a correlated metallic state for $x > 0.04$. In order to reveal the nature of the metallic state, a detailed study of the elementary excitations is required that can determine the doping evolution of the electronic interactions, especially the magnetic couplings. Furthermore, since the magnetic anisotropy has been suggested to contain bond-directional pseudodipolar interactions arising from the $J_{\text{eff}} = \frac{1}{2}$ states, its response to

electron doping will shed light on understanding the SOC physics in doped iridates [5,8,29,30].

In this Letter, we present measurements of the doping dependence of the magnetic order and the elementary excitations across the electron-doping driven IMT in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ ($x = 0, 0.02, 0.03, 0.05$, and 0.065) using Ir- L_3 edge resonant inelastic x-ray scattering (RIXS) [36]. Our results reveal an evolution of the AFM from three-dimensional (3D) long range AFM (LAF) ($x \leq 0.03$) to 3D short range AFM (SAF) ($x \sim 0.05$) across the IMT and subsequent 2D SAF deep in the metallic state ($x = 0.065$). Following the evolution of the magnetic order, we present a detailed analysis of the doping-dependent magnetic excitations and spin-orbit excitons, from which the doping effects on the magnetic couplings and the nature of the metallic state are determined.

The measurements on $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ were carried out at the ID20 beam line ($x = 0.02, 0.05$, and 0.065) of the European Synchrotron Radiation Facility (ESRF) and the 27ID-B beam line ($x = 0$ and 0.03) at the Advanced Photon Source (APS) [37–39]. To facilitate comparison with previous results [29,30], we use the tetragonal notation [Figs. 1(a) and 1(b)] in presenting our RIXS results and define the momentum transfer \mathbf{Q} in reciprocal space as $\mathbf{Q} = H\mathbf{a}^* + K\mathbf{b}^* + L\mathbf{c}^*$, where H , K , and L are Miller indices and $\mathbf{a}^* = \hat{\mathbf{a}}(2\pi/a)$, $\mathbf{b}^* = \hat{\mathbf{b}}(2\pi/b)$, $\mathbf{c}^* = \hat{\mathbf{c}}(2\pi/c)$ with $a \approx b \approx 3.9$ Å, and $c \approx 20.9$ Å for $\text{Sr}_3\text{Ir}_2\text{O}_7$. In this notation, the wave vector of the c -axis G -type AFM is $\mathbf{q} = (\frac{1}{2}, \frac{1}{2}, 0)$ [29,30].

We first describe the doping evolution of the magnetic order. Figure 1(a) shows the in-plane structure and the rotations of the IrO_6 octahedra in $\text{Sr}_3\text{Ir}_2\text{O}_7$. The exchange couplings on the tetragonal lattice and their naming conventions are described in Fig. 1(b). Figure 1(c) is a schematic magnetic and electronic phase diagram of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ [31,32]. The doping levels x measured at $T = 30$ K are indicated by red circles. To characterize the doping-dependent AFM, we have measured the magnetic Bragg peaks along $[H, \frac{1}{2}, 28]$ and $[\frac{1}{2}, \frac{1}{2}, L]$ for $x = 0, 0.02, 0.03, 0.05$, and 0.065 using the elastic channel of the RIXS spectrometer [39]. The L scan for $x = 0$ displays magnetic Bragg peaks from $L = 19$ to 28 with an intensity modulation [Fig. 1(d)], which has a period controlled by the ratio between lattice parameter c and bilayer distance (d) ($(c/d) \approx 5.1$) [8]. Upon electron doping, the 3D LAF persists for $x = 0.03$ but becomes short ranged for $x = 0.05$, as indicated by the broad peaks along both H and L in Figs. 1(e) and 1(f). The L scan for the metallic $x = 0.05$ sample deserves special attention. It reveals a broad feature superimposed on a flat background, and is well fitted by a sum of the bilayer antiferromagnetic structural factor $\cos^2[(2\pi d/c)]$ [8] and a constant background [Fig. 1(f)]. This broad feature indicates that the magnetic correlation length along the c axis has decreased to a very small value comparable with the bilayer distance.

This suggests that the c -axis magnetic correlations supporting the G -type AFM are about to disappear. The constant background can be attributed to a vanishing of the 3D SAF in a partial volume of the sample. This is in agreement with the percolative nature of the IMT [32]. For $x = 0.065$, the L scan becomes featureless while the broader in-plane magnetic Bragg peak remains and persists at 290 K [Figs. 1(e) and 1(f)]. This indicates that further doping suppresses the magnetic couplings that support the 3D SAF and drives the system into a robust 2D SAF state.

To understand the doping-dependent electronic interactions across the IMT and the transitions between 3D LAF and 2D SAF, we have presented the magnetic excitations and the spin-orbit excitons of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ in Figs. 2

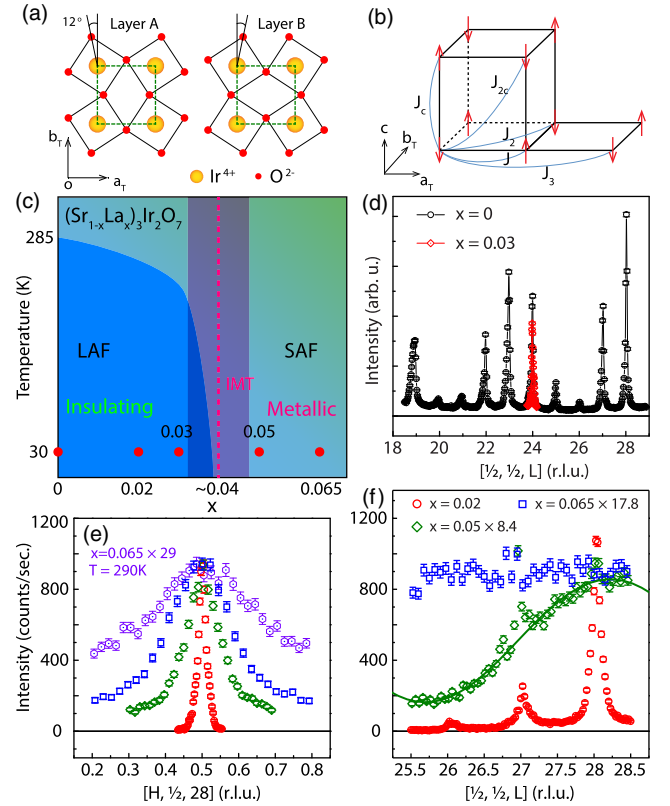


FIG. 1. (a) In-plane crystal structure and the rotation of the IrO_6 octahedra ($\sim 12^\circ$) around the c axis. (b) G -type collinear antiferromagnetic order with moment along the c axis. J_1, J_2 , and J_3 are first, second, and third nearest superexchange couplings within the ab plane, respectively. J_c and J_{2c} are first and second nearest couplings along the c axis. (c) Schematic phase diagram of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$, adapted from Refs. [31,32]. LAF and SAF are long range and short range antiferromagnetic order, respectively. The insulator-to-metal transition (IMT) occurs at $x \sim 0.04$. The dopings used in the present study are marked by red dots. (d) L scan of the c -axis G -type antiferromagnetic order for $x = 0$ and 0.03 . (e), (f) Doping dependent H and L scans across the magnetic Bragg peak $(\frac{1}{2}, \frac{1}{2}, 28)$. The green solid curve in (e) is a fit of the L scan for $x = 0.05$. All the measurements were performed at 30 K unless otherwise indicated.

and 3. The in-plane momentum dependent RIXS for $x = 0$, 0.02 and 0.065 are shown as color maps in Figs. 2(a)–2(c). For $x = 0$, the dispersion, the large magnon gap, and the spectral-weight distribution are consistent with a previous report measured at the same L [29]. In addition, our results reveal clear dispersive spin-orbit excitons exhibiting similar energy scale and dispersion to that in Sr_2IrO_4 [7,11,12,50,51]. As increasing x , the magnetic excitations are damped: they broaden in energy and decrease in intensity. This damping has been reported in other charge-carrier doped 2D correlated systems such as cuprates and iron pnictides [52–56]. The spin-orbit excitons show weak doping dependence, indicating that the magnetic excitations are fluctuations of the robust $J_{\text{eff}} = \frac{1}{2}$ pseudospins [39].

The strong doping dependence of the magnetic excitations are depicted in Figs. 2 and 3. The dispersions are symmetric about $(\frac{1}{4}, \frac{1}{4})$ and change less from $x = 0$ to $x = 0.03$ [Fig. 3(a)]. Across the IMT to $x = 0.05$, the dispersion becomes asymmetric and exhibits different gap sizes at $(\frac{1}{2}, \frac{1}{2})$ and $(0, 0)$. A substantial softening occurs along $(\frac{1}{2}, \frac{1}{2})$ – $(\frac{1}{4}, \frac{1}{4})$ while the band top at $(\frac{1}{2}, 0)$ remains unchanged [Figs. 2 and 3(b)]. This anisotropic softening is followed by a further softening at $(\frac{1}{4}, \frac{1}{4})$ and, surprisingly, a sizable hardening at $(\frac{1}{2}, 0)$ in $x = 0.065$ [Figs. 2 and 3]. Furthermore, the large magnon gap for $x \leq 0.05$ collapses dramatically in $x = 0.065$ [Figs. 2(f), 2(g), and 3(b)], where the magnetic excitations at $(\frac{1}{2}, \frac{1}{2})$ overlap with the elastic

magnetic scattering [Fig. 2(f)], whereas a weak signal is observed at $(0, 0)$ [Fig. 2(g)] [39]. A similar anisotropic softening between $(\frac{1}{4}, \frac{1}{4})$ and $(\frac{1}{2}, 0)$ was observed in $\text{Sr}_{2-x}\text{La}_x\text{IrO}_4$ and attributed to the interaction between magnetic moments and emergent itinerant electrons having developed a Fermi surface at $(\frac{1}{4}, \frac{1}{4})$ [22–24]. The same explanation can be applied here since La introduces itinerant electrons and Fermi pockets have been well developed at $(\frac{1}{4}, \frac{1}{4})$ in metallic $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ [33–35]. Therefore the anisotropic softening is in line with that in $\text{Sr}_{2-x}\text{La}_x\text{IrO}_4$, indicating a similar role of the emergent itinerant electrons.

The hardening of the magnetic excitations at $(\frac{1}{2}, 0)$ and the collapse of the magnon gap occur between $x = 0.05$ and 0.065 where the system evolves from 3D SAF to 2D SAF. Since the AFM along the c axis is absent in 2D SAF, we expect that the interlayer couplings will be greatly suppressed in $x = 0.065$. With the suppression of the interlayer couplings, $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ could be similar to the single-layer $\text{Sr}_{2-x}\text{La}_x\text{IrO}_4$ ($x \geq 0.04$), in which the magnetic excitations are gapless and have a larger band top at $(\frac{1}{2}, 0)$ [23]. Indeed, we find the 2D pseudospin- $\frac{1}{2}$ ($J - J_2 - J_3$) model used for Sr_2IrO_4 describes the dispersion and gives similar fitting parameters ($J = 44, J_2 = -29, J_3 = 14.4$ meV) to $\text{Sr}_{2-x}\text{La}_x\text{IrO}_4$ ($x \geq 0.04$) [Fig. 3], indicating that electron doping drives the bilayer iridate $\text{Sr}_3\text{Ir}_2\text{O}_7$ into a 2D magnetic system exhibiting strong antiferromagnetic pseudospin

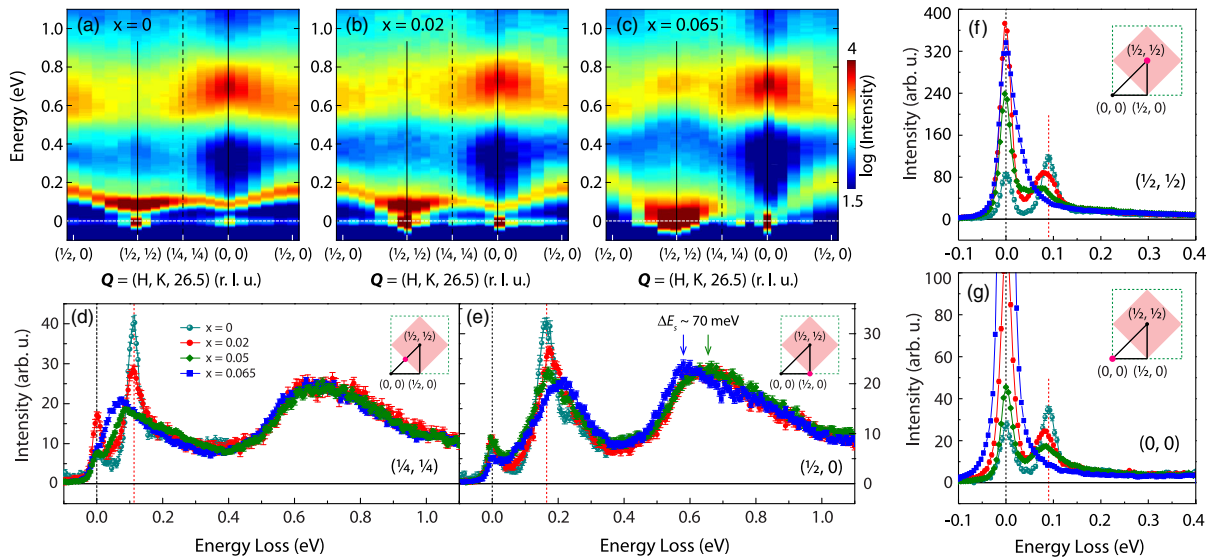


FIG. 2. (a)–(c) In-plane momentum dependence of RIXS spectra of $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ for $x = 0, 0.02$, and 0.065 . (d), (e) Comparison of the elementary excitations between $x = 0, 0.02, 0.05$, and 0.065 collected at $Q = (\frac{1}{4}, \frac{1}{4})$ and $(\frac{1}{2}, 0)$. For $x = 0$, only magnetic excitations are shown. (f), (g) Doping-dependent magnetic excitations at $Q = (\frac{1}{2}, \frac{1}{2})$ and $(0, 0)$. The insets in (d)–(g) illustrate the reciprocal space where the green dashed square and the pink solid square are the tetragonal Brillouin zone and the antiferromagnetic Brillouin zone, respectively. The pink filled circles mark the vector positions for (d)–(g). The vertical red dashed lines mark the peak position of the magnons for $x = 0$. The blue and green arrows in (e) show the peak positions of the spin-orbit excitons and ΔE_s marks the energy difference of the peak positions.

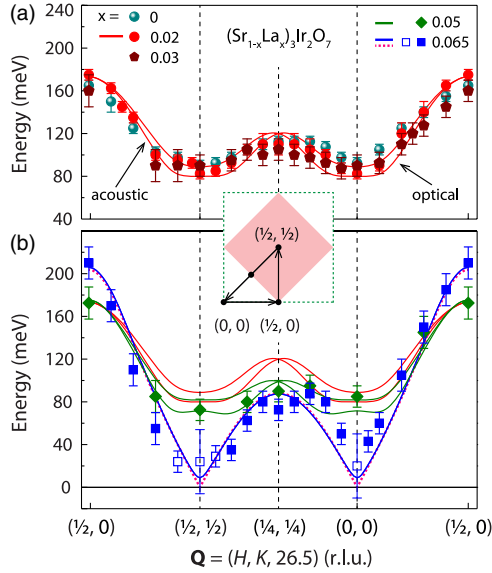


FIG. 3. Doping-dependent magnon dispersions for $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$. To show the data clearly, the dispersions are split into two panels (a) $x = 0, 0.02$, and 0.03 and (b) $x = 0.05$ and 0.065 . The dispersions for $x \leq 0.05$ and the blue solid squares of $x = 0.065$ are obtained by selecting the peak positions of the magnetic excitations. The blue open squares of $x = 0.065$ are extracted from fitting of the magnetic excitations [39]. The solid curves are fits to the dispersions for $x = 0.02, 0.05$, and 0.065 using the bilayer model, which includes an acoustic and an optical branch [29]. The pink dashed curve is the fitting of the dispersion for 0.065 using the $J - J_2 - J_3$ model [23].

fluctuations [23]. The strong magnetic excitations in $x = 0.065$ demonstrate that metallic $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ hosts strong electron correlations like its parent compound, and therefore provide a solid evidence for a correlated metallic picture [32].

To quantitatively understand the doping evolution of the magnetic couplings, we have fitted the dispersions using the bilayer model [Fig. 3] [29,39], which consists of nearest neighbor (H_0) and long range interactions (H_1)

$$H_0 = \sum_{\langle i,j \rangle} [J_{ij} \vec{S}_i \cdot \vec{S}_j + \Gamma_{ij} S_i^z S_j^z + \vec{D}_{ij} \cdot (\vec{S}_i \times \vec{S}_j)] \quad (1)$$

$$H_1 = \sum_{\langle\langle i,j \rangle\rangle} J_2 \vec{S}_i \cdot \vec{S}_j + \sum_{\langle\langle\langle i,j \rangle\rangle\rangle} J_3 \vec{S}_i \cdot \vec{S}_j + \sum_{\langle i,j \rangle} J_{2c} \vec{S}_i \cdot \vec{S}_{j+z} \quad (2)$$

where Γ_{ij} is the anisotropic coupling including the bond-directional pseudodipolar interactions and D_{ij} the Dzyaloshinsky-Moriya interaction arising from the staggered rotation of the IrO_6 octahedra [Fig. 1(a)]. H_0 contains the sum over both intralayer ($J_{ij} = J$, $\Gamma_{ij} = \Gamma$, and $D_{ij} = D$) and interlayer couplings (J_c , Γ_c , and D_c). The $\langle i, j \rangle$, $\langle\langle i, j \rangle\rangle$, and $\langle\langle\langle i, j \rangle\rangle\rangle$ in H_1 represent in-plane first, second, and third nearest neighbors and J_2 , J_3 , and J_{2c} the long range exchange couplings [Fig. 1(b)]. The c -axis couplings J_c and J_{2c} and D_c are responsible for the bilayer

splitting of the acoustic and optical branches, and the magnon gap arises from the anisotropic couplings Γ and Γ_c [29]. Because of the presence of interlayer couplings, our fittings for $x \leq 0.05$ have been restricted to adjusting only J , J_2 , and J_3 [23,39]. For $x = 0.065$, we fit all the parameters to account for the suppression of the interlayer couplings and the magnetic anisotropy. As shown in Fig. 3, our fitting successfully describes the anisotropic softening and the collapse of the magnon gap [39]. With increasing doping, the anisotropic softening is captured by the evolution of J and J_2 [39], indicative of strong interplay between the in-plane nearest and next nearest couplings and the emergent itinerant electrons. The bilayer splitting of the two branches disappear in 2D SAF state since J_c , J_{2c} , and D_c vanish. The collapse of the magnon gap corresponds to the suppression of the anisotropic couplings (Γ and Γ_c) including the bond-directional pseudodipolar interactions [39]. With the suppression of the interlayer couplings and the magnetic anisotropy, the bilayer model is similar to the $J - J_2 - J_3$ model, as indicated by the fitting curves using these two models [Fig. 3(b)].

Although the model describes the anisotropic softening and gap collapse, it fails to capture the asymmetry of the dispersions for metallic $x = 0.05$ and 0.065 since it is intrinsically symmetric about $(\frac{1}{4}, \frac{1}{4})$. As shown in Fig. 3(b), the dispersion lies below the fitting on the left of $(\frac{1}{4}, \frac{1}{4})$ but above the fitting on the right side. We attribute this to different damping rates between $(\frac{1}{2}, \frac{1}{2})$ and $(0,0)$ driven by certain interactions between the pseudospins and the emergent itinerant electrons, which are not taken into account by the bilayer model developed for the insulating antiferromagnetic $\text{Sr}_3\text{Ir}_2\text{O}_7$ [5,8,29]. Therefore, understanding the microscopic mechanism for the interactions requires further studies. Nonetheless, the effective fitting partially reflects the doping effects on the magnetic interactions and the dynamics of the pseudospins in the context of the emergent itinerant electrons. We have also tried to describe the dispersions using the quantum-dimer model reported in Ref. [30]. The results are presented in the Supplemental Material [39].

We now turn to the electron-doping effects on the spin-orbit excitons in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$. Besides demonstrating the robustness of the $J_{\text{eff}} = \frac{1}{2}$ moments, the spin-orbit excitons undergo a sudden decrease in energy ($\Delta E_s \sim 70$ meV) at momentum close to $(\frac{1}{2}, 0)$ between $x = 0.05$ and 0.065 . Since the spin-orbit excitons are electronic transitions from the $j = \frac{3}{2}$ band to the $j = \frac{1}{2}$ upper Hubbard band, this energy decrease suggests that these two bands are modified across the 3D-to-2D SAF transition [39]. This implies that the changes of magnetic order and magnetic couplings could induce band renormalization in this system [39].

In summary, our measurements on $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ unveil an electron-doping evolution of the AFM from

3D LAF to 3D SAF and subsequent 2D SAF deep in the metallic state and illustrate a more detailed phase diagram [31,32]. We show subsequently that the magnons undergo an anisotropic damping with increasing doping, with the large magnon gap strongly suppressed in the 2D SAF metallic regime. This indicates that the emergent itinerant electrons suppress the AFM by weakening the magnetic couplings and drive the system into a 2D SAF correlated metallic state hosting strong antiferromagnetic fluctuations of the $J_{\text{eff}} = \frac{1}{2}$ moments. Our results provide a solid experimental basis that will guide future theoretical works on the physics of doping the SOC-induced Mott insulators in the presence of strong interlayer couplings.

We thank Matteo Rossi (ESRF) for helpful discussions. The work at PSI is supported by the Swiss National Science Foundation through its Sinergia network Mott Physics Beyond the Heisenberg Model (MPBH) and the NCCR-MARVEL. Xingye Lu acknowledges financial support from the European Community's Seventh Framework Programme (FP7/2007–2013) under Grant agreement No. 290605 (COFUND: PSI-FELLOW). G. Cao acknowledges support by the US National Science Foundation via Grants No. DMR-1265162 and No. DMR-1712101. The work used Sector 27 of the Advanced Photon Source, a U.S. Department of Energy (DOE) Office of Science User Facility operated for the DOE Office of Science by Argonne National Laboratory under Contract No. DE-AC02-06CH11357.

Note added.—Asymmetric dispersion of the magnetic excitations have recently also been observed by Hogan *et al.* in $(\text{Sr}_{1-x}\text{La}_x)_3\text{Ir}_2\text{O}_7$ [57], where the dispersion for $x = 0.07$ is much akin to our results on $x = 0.05$.

*xingye.lu@psi.ch

†thorsten.schmitt@psi.ch

- [1] W. Witczak-Krempa, G. Chen, Y. B. Kim, and L. Balents, *Annu. Rev. Condens. Matter Phys.* **5**, 57 (2014).
- [2] D. Pesin and L. Balents, *Nat. Phys.* **6**, 376 (2010).
- [3] J. G. Rau, E. Kin-Ho Lee, and H.-Y. Kee, *Annu. Rev. Condens. Matter Phys.* **7**, 195 (2016).
- [4] B. J. Kim, H. Jin, S. J. Moon, J.-Y. Kim, B.-G. Park, C. S. Leem, J. Yu, T. W. Noh, C. Kim, S.-J. Oh, J.-H. Park, V. Durairaj, G. Cao, and E. Rotenberg, *Phys. Rev. Lett.* **101**, 076402 (2008).
- [5] G. Jackeli and G. Khaliullin, *Phys. Rev. Lett.* **102**, 017205 (2009).
- [6] B. J. Kim, H. Ohsumi, T. Komesu, S. Sakai, T. Morita, H. Takagi, and T. Arima, *Science* **323**, 1329 (2009).
- [7] S. J. Moon, H. Jin, K. W. Kim, W. S. Choi, Y. S. Lee, J. Yu, G. Cao, A. Sumi, H. Funakubo, C. Bernhard, and T. W. Noh, *Phys. Rev. Lett.* **101**, 226402 (2008).
- [8] J. W. Kim, Y. Choi, Jungho Kim, J. F. Mitchell, G. Jackeli, M. Daghofer, J. van den Brink, G. Khaliullin, and B. J. Kim, *Phys. Rev. Lett.* **109**, 037204 (2012).
- [9] P. A. Lee, N. Nagaosa, and X.-G. Wen, *Rev. Mod. Phys.* **78**, 17 (2006).
- [10] F. Ye, S. Chi, B. C. Chakoumakos, J. A. Fernandez-Baca, T. Qi, and G. Cao, *Phys. Rev. B* **87**, 140406 (2013).
- [11] J. Kim, D. Casa, M. H. Upton, T. Gog, Y.-J. Kim, J. F. Mitchell, M. van Veenendaal, M. Daghofer, J. van den Brink, G. Khaliullin, and B. J. Kim, *Phys. Rev. Lett.* **108**, 177003 (2012).
- [12] J. Kim, M. Daghofer, A. H. Said, T. Gog, J. van den Brink, G. Khaliullin, and B. J. Kim, *Nat. Commun.* **5**, 4453 (2014).
- [13] R. Coldea, S. M. Hayden, G. Aeppli, T. G. Perring, C. D. Frost, T. E. Mason, S.-W. Cheong, and Z. Fisk, *Phys. Rev. Lett.* **86**, 5377 (2001).
- [14] N. S. Headings, S. M. Hayden, R. Coldea, and T. G. Perring, *Phys. Rev. Lett.* **105**, 247001 (2010).
- [15] F. Wang and T. Senthil, *Phys. Rev. Lett.* **106**, 136402 (2011).
- [16] H. Watanabe, T. Shirakawa, and S. Yunoki, *Phys. Rev. Lett.* **110**, 027002 (2013).
- [17] M. Ge, T. F. Qi, O. B. Korneta, D. E. De Long, P. Schlottmann, W. P. Crummett, and G. Cao, *Phys. Rev. B* **84**, 100402(R) (2011).
- [18] X. Chen, T. Hogan, D. Walkup, W. Zhou, M. Pokharel, M. Yao, W. Tian, T. Z. Ward, Y. Zhao, D. Parshall, C. Opeil, J. W. Lynn, V. Madhavan, and S. D. Wilson, *Phys. Rev. B* **92**, 075125 (2015).
- [19] Y. K. Kim, O. Krupin, J. D. Denlinger, A. Bostwick, E. Rotenberg, Q. Zhao, J. F. Mitchell, J. W. Allen, and B. J. Kim, *Science* **345**, 187 (2014).
- [20] Y. J. Yan, M. Q. Ren, H. C. Xu, B. P. Xie, R. Tao, H. Y. Choi, N. Lee, Y. J. Choi, T. Zhang, and D. L. Feng, *Phys. Rev. X* **5**, 041018 (2015).
- [21] Y. K. Kim, N. H. Sung, J. D. Denlinger, and B. J. Kim, *Nat. Phys.* **12**, 37 (2016).
- [22] A. de la Torre, S. McKeown Walker, F. Y. Bruno, S. Ricco, Z. Wang, I. Gutierrez Lezama, G. Scheerer, G. Giriat, D. Jaccard, C. Berthod, T. K. Kim, M. Hoesch, E. C. Hunter, R. S. Perry, A. Tama, and F. Baumberger, *Phys. Rev. Lett.* **115**, 176402 (2015).
- [23] H. Gretarsson, N. H. Sung, J. Porras, J. Bertinshaw, C. Dietl, J. A. N. Bruin, A. F. Bangura, Y. K. Kim, R. Dinnebier, J. Kim, A. Al-Zein, M. Moretti Sala, M. Krisch, M. Le Tacon, B. Keimer, and B. J. Kim, *Phys. Rev. Lett.* **117**, 107001 (2016).
- [24] X. Liu, M. P. M. Dean, Z. Y. Meng, M. H. Upton, T. Qi, T. Gog, Y. Cao, J. Q. Lin, D. Meyers, H. Ding, G. Cao, and J. P. Hill, *Phys. Rev. B* **93**, 241102(R) (2016).
- [25] D. Reznik, P. Bourges, H. F. Fong, L. P. Regnault, J. Bossy, C. Vettier, D. L. Milius, I. A. Aksay, and B. Keimer, *Phys. Rev. B* **53**, R14741(R) (1996).
- [26] J. Nichols, N. Bray-Ali, A. Ansary, G. Cao, and Kwok-Wai Ng, *Phys. Rev. B* **89**, 085125 (2014).
- [27] Y. Okada, D. Walkup, H. Lin, C. Dhital, T.-R. Chang, S. Khadka, W. Zhou, H.-T. Jeng, M. Paranjape, A. Bansil, Z. Wang, S. D. Wilson, and V. Madhavan, *Nat. Mater.* **12**, 707 (2013).
- [28] S. Boseggia, R. Springell, H. C. Walker, A. T. Boothroyd, D. Prabhakaran, D. Wermeille, L. Bouchenoire, S. P. Collins, and D. F. McMorrow, *Phys. Rev. B* **85**, 184432 (2012).

- [29] J. Kim, A. H. Said, D. Casa, M. H. Upton, T. Gog, M. Daghofer, G. Jackeli, J. van den Brink, G. Khaliullin, and B. J. Kim, *Phys. Rev. Lett.* **109**, 157402 (2012).
- [30] M. Moretti Sala, V. Schnells, S. Boseggia, L. Simonelli, A. Al-Zein, J. G. Vale, L. Paolasini, E. C. Hunter, R. S. Perry, D. Prabhakaran, A. T. Boothroyd, M. Krisch, G. Monaco, H. M. Ronnow, D. F. McMorro, and F. Mila, *Phys. Rev. B* **92**, 024405 (2015).
- [31] L. Li, P. P. Kong, T. F. Qi, C. Q. Jin, S. J. Yuan, L. E. DeLong, P. Schlottmann, and G. Cao, *Phys. Rev. B* **87**, 235127 (2013).
- [32] T. Hogan, Z. Yamani, D. Walkup, X. Chen, R. Dally, T. Z. Ward, M. P. M. Dean, J. Hill, Z. Islam, V. Madhavan, and S. D. Wilson, *Phys. Rev. Lett.* **114**, 257203 (2015).
- [33] A. de la Torre, E. C. Hunter, A. Subedi, S. McKeown Walker, A. Tamai, T. K. Kim, M. Hoesch, R. S. Perry, A. Georges, and F. Baumberger, *Phys. Rev. Lett.* **113**, 256402 (2014).
- [34] J. He, H. Hafiz, T. R. Mion, T. Hogan, C. Dhital, X. Chen, Q. Lin, M. Hashimoto, D. H. Lu, Y. Zhang, R. S. Markiewicz, A. Bansil, S. D. Wilson, and R.-H. He, *Sci. Rep.* **5**, 8533 (2015).
- [35] J. He *et al.*, *Nat. Mater.* **14**, 577 (2015).
- [36] L. J. P. Ament, M. van Veenendaal, T. P. Devereaux, J. P. Hill, and J. van den Brink, *Rev. Mod. Phys.* **83**, 705 (2011).
- [37] M. Moretti Sala, C. Henriquet, L. Simonelli, R. Verbeni, and G. Monaco, *J. Electron Spectrosc. Relat. Phenom.* **188**, 150 (2013).
- [38] Yu. V. Shvyd'ko, J. P. Hill, C. A. Burns, D. S. Coburn, B. Brajuskovic, D. Casa, K. Goetze, T. Gog, R. Khachatryan, J.-H. Kim, C. N. Koditwaku, M. Ramanathan, T. Roberts, A. Said, H. Sinn, D. Shu, S. Stoupin, M. Upton, M. Wiczorek, and H. Yavas, *J. Electron Spectrosc. Relat. Phenom.* **188**, 140 (2013).
- [39] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevLett.118.027202> for details, which includes Refs. [40–49].
- [40] G. Cao, Y. Xin, C. S. Alexander, J. E. Crow, P. Schlottmann, M. K. Crawford, R. L. Harlow, and W. Marshall, *Phys. Rev. B* **66**, 214412 (2002).
- [41] H. Gretarsson, N. H. Sung, M. Höppner, B. J. Kim, B. Keimer, and M. Le Tacon, *Phys. Rev. Lett.* **116**, 136401 (2016).
- [42] L. W. Harriger, M. S. Liu, H. Q. Luo, R. A. Ewings, C. D. Frost, T. G. Perring, and P. Dai, *Phys. Rev. B* **86**, 140403 (2012).
- [43] H. Zhang, K. Haule, and D. Vanderbilt, *Phys. Rev. Lett.* **111**, 246402 (2013).
- [44] C. Monney, K. J. Zhou, H. Cercellier, Z. Vydrova, M. G. Garnier, G. Monney, V. N. Strocov, H. Berger, H. Beck, T. Schmitt, and P. Aebi, *Phys. Rev. Lett.* **109**, 047401 (2012).
- [45] R. Arita, J. Kuneš, A. V. Kozhevnikov, A. G. Eguiluz, and M. Imada, *Phys. Rev. Lett.* **108**, 086403 (2012).
- [46] D. Hsieh, F. Mahmood, D. H. Torchinsky, G. Cao, and N. Gedik, *Phys. Rev. B* **86**, 035128 (2012).
- [47] Q. Li, G. Cao, S. Okamoto, J. Yi, W. Lin, B. C. Sales, J. Yan, R. Arita, J. Kuneš, A. V. Kozhevnikov, A. G. Eguiluz, M. Imada, Z. Gai, M. Pan, and D. G. Mandrus, *Sci. Rep.* **3**, 3073 (2013).
- [48] J. Matsuno, K. Ihara, S. Yamamura, H. Wadati, K. Ishii, V. V. Shankar, H. Y. Kee, and H. Takagi, *Phys. Rev. Lett.* **114**, 247209 (2015).
- [49] H. Zheng, J. Terzic, Feng Ye, X. G. Wan, D. Wang, Jinchun Wang, Xiaoping Wang, P. Schlottmann, S. J. Yuan, and G. Cao, *Phys. Rev. B* **93**, 235157 (2016).
- [50] Y. Ding, L. Yang, C. C. Chen, H. S. Kim, M. J. Han, W. Luo, Z. Feng, M. Upton, D. Casa, Jungho Kim, T. Gog, Z. Zeng, G. Cao, H. K. Mao, and M. van Veenendaal, *Phys. Rev. Lett.* **116**, 216402 (2016).
- [51] S. Bossegia, Ph.D. thesis, University College London, 2014, pp. 182, 184, 193.
- [52] M. Le Tacon, G. Ghiringhelli, J. Chaloupka, M. Moretti Sala, V. Hinkov, M. W. Haverkort, M. Minola, M. Bakr, K. J. Zhou, S. Blanco-Canosa, C. Monney, Y. T. Song, G. L. Sun, C. T. Lin, G. M. De Luca, M. Salluzzo, G. Khaliullin, T. Schmitt, L. Braicovich, and B. Keimer, *Nat. Phys.* **7**, 725 (2011).
- [53] M. P. M. Dean, G. Dellea, R. S. Springell, F. Yakhou-Harris, K. Kummer, N. B. Brookes, X. Liu, Y.-J. Sun, J. Strle, T. Schmitt, L. Braicovich, G. Ghiringhelli, I. Bozovic, and J. P. Hill, *Nat. Mater.* **12**, 1019 (2013).
- [54] K.-J. Zhou, Y.-B. Huang, C. Monney, X. Dai, V. N. Strocov, N.-L. Wang, Z.-G. Chen, C. Zhang, P. Dai, L. Patthey, J. van den Brink, H. Ding, and T. Schmitt, *Nat. Commun.* **4**, 1470 (2013).
- [55] J. Pelliciari, Y. Huang, T. Das, M. Dantz, V. Bisogni, P. O. Velasco, V. N. Strocov, L. Xing, X. Wang, C. Jin, and T. Schmitt, *Phys. Rev. B* **93**, 134515 (2016).
- [56] P. Dai, *Rev. Mod. Phys.* **87**, 855 (2015).
- [57] T. Hogan, R. Dally, M. Upton, J. P. Clancy, K. Finkelstein, Y.-J. Kim, M. J. Graf, and S. D. Wilson, *Phys. Rev. B* **94**, 100401(R) (2016).